

Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/GB05/001159

International filing date: 24 March 2005 (24.03.2005)

Document type: Certified copy of priority document

Document details: Country/Office: GB
Number: 0406841.7
Filing date: 26 March 2004 (26.03.2004)

Date of receipt at the International Bureau: 26 May 2005 (26.05.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse



INVESTOR IN PEOPLE

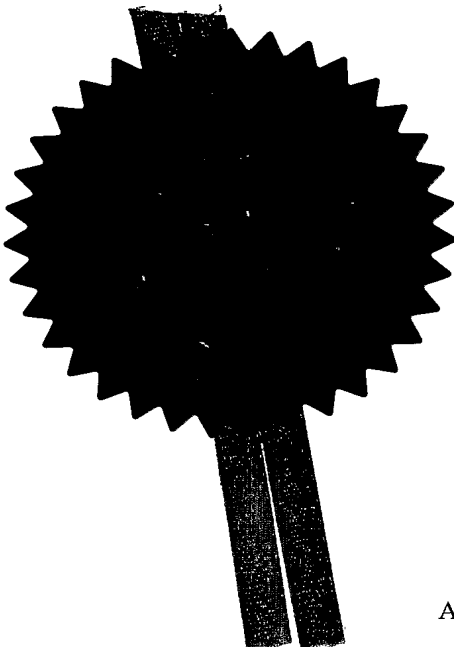
The Patent Office
Concept House
Cardiff Road
Newport
South Wales
NP10 8QQ

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

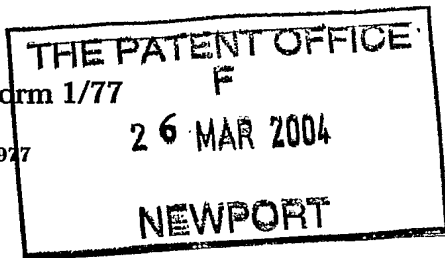


Signed

Andrew Gersey

Dated 22 April 2005





974222*97#*977Y?#?DS?
29MAR04 E884252-1 D00239
P01/7700 0.00-0406841.7 NONE

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

26 MAR 2004

The Patent Office

Cardiff Road
Newport
South Wales
NP10 8QQ

1. Your reference JTS\IP13212GB

2. Patent application number
(The Patent Office will fill in this part)

0406841.7

3. Full name, address and postcode of the or of each applicant (underline all surnames)

The University Court of the University of St. Andrews
College Gate
North Street
St. Andrews
Fife, KY16 9AJ, Scotland, UK

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

06512446001

4. Title of the invention
Surface Patterning System

5. Name of your agent (if you have one)

Cruikshank & Fairweather

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

19 Royal Exchange Square
Glasgow, G1 3AE
Scotland, UK

Patents ADP number (if you know it)

547002 ✓

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)Date of filing
(day / month / year)

-

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

-

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

- a) any applicant named in part 3 is not an inventor, or
b) there is an inventor who is not named as an applicant, or
c) any named applicant is a corporate body.
See note (d))

Yes

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form.
Do not count copies of the same document

Continuation sheets of this form

Description 17 -

Claim(s) -

Abstract -

Drawing(s) 3 13 02

10. If you are also filing any of the following, state how many against each item.

Priority documents -

Translations of priority documents -

Statement of inventorship and right to grant of a patent (Patents Form 7/77) -

Request for preliminary examination and search (Patents Form 9/77) -

Request for substantive examination (Patents Form 10/77) -

Any other documents (please specify) -

11. I/We request the grant of a patent on the basis of this application.

Signature

Jan Szczuka

Date

25/03/04

12. Name and daytime telephone number of person to contact in the United Kingdom Mr Jan Szczuka - 0131 225 4500

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 08459 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

PATTERNING METHODS AND PRODUCTS

The present invention relates to methods and processes of patterning self-assembled mono-layers, and product obtainable by such processes.

5

There is considerable interest in the production of SAMs for various purposes. Recently there has been disclosed in WO 0123962 a surface-modified layer system in which a self-assembled monolayer (SAM) has been irradiated by various kinds of radiation selected from electron beam, plasma, X-Ray, β -Ray, γ -Ray and UV which results in cross-linking of the molecules of the SAM thereby forming a protective coating on the substrate which is resistant to damage caused by friction or corrosion.

10

15

The use of such types of highly energetic radiation which result in chemical change inevitably involve a degree of lack of control of the chemical reactions involved and can result in the incidence of undesired side-reactions, and/or the production of undesirable artefacts and by-products. Furthermore,

20

electrochemical and electronic properties are inevitably changed upon irradiation of SAMs with energetic particles rendering for example, such SAMs electrically non-conducting or chemically inert. Another problem is that, in the case of radiation in the form of particles such as electrons, ions, plasma, special

25

environments (e.g. vacuum) are required.

There is a need for improved and/or alternative techniques for production of stable SAMs, especially patterned SAMs. It is also an object of the invention to avoid or minimise one or more disadvantages of the prior art.

5

It has now been found that by selection of the compounds used to form the SAM, it is possible to obtain a polymorphic SAM which can be transformed from a first, less stable, structure, to a second, more stable, structure, by means of thermal treatment thereof. In more detail, it has been found that polymorphic SAMs can be obtained by using selected aryl moiety species in which at least two of the various factors affecting the energy balance in the SAM on the particular substrate used, are in competition with each other - in contrast to the normal practice employed in producing SAMs, where, in some cases, the compounds used in the SAMs are selected so as to minimise energy with all the various factors affecting the energy balance being used in collaboration with each other to maximise the stability of the SAM, or in other cases, no attention at all is paid to how different factors affect the energy balance. Furthermore it has been found that the, less stable, SAMs produced with such selected compounds in accordance with the present invention, can be thermally patterned to transfer an image to the SAM, in which the image is defined by areas with different structural forms. Still further it has been found that these different structural forms have different degrees of stability whereby such a

10

15

20

25

"latent" image - defined "only" by differences in structural form, can be "developed" or "fixed" by means of suitable processing of one or other of the thermally treated and untreated parts of the SAM, for example, by exchange, or
5 otherwise removal, of the less stable structural form-SAM, thereby resulting in a chemically, and/or structurally, and/or topographically defined pattern.

In one aspect the present invention provides a surface-modified
10 layer system comprising a substrate having a surface and a self-assembled monolayer (SAM) anchored to at least part of said surface, wherein said SAM is comprised by an aryl moiety species in a substantially stable structural form derived, in situ, by thermal treatment from a less stable structural form.

15

The invention also provides a process for producing a surface-modified layer system comprising a substrate having a surface and a self-assembled mono-layer (SAM) anchored to at least part of said surface, wherein said SAM is comprised by an aryl moiety
20 species, said surface comprising the steps of:

- a) providing a SAM anchored to a substrate, wherein said SAM is polymorphic having at least first and second structural forms; and
- b) thermally treating said SAM so as to change said SAM from
25 said first structural form to said second structural form.

In another aspect the present invention provides a thermo-lithographic process comprising the steps of:

- 5 a) providing a SAM anchored to a substrate, wherein said SAM is polymorphic having at least first and second structural forms; and
- b) transferring a desired pattern to said SAM using thermal treatment so as to change selectively part of said SAM from said first structural form to said second structural form.

10

In general said process also includes the further step of developing the thermally treated SAM by subjecting it to further processing so as to substantially modify selectively one of: thermally treated and non-thermally treated parts of the SAM.

15

In another aspect the invention provides a method of providing a surface-modified layer system comprising a SAM anchored to a substrate surface in a desired pattern thereon, which method comprises the steps of:

- 20 a) providing a substrate and a compound having a selected aryl moiety species and an anchor moiety bondable to said substrate so as to exert a directive force with respect to the molecular orientation, said selected aryl moiety species having a plurality of different parameters affecting the energy-balance
- 25 of the SAM, wherein at least two of said parameters exert opposing directive forces arising from the molecule-substrate

interaction which have a substantially competitive effect on said energy-balance which results in polymorphism of said SAM;

b) bonding of said compound to said substrate; and

c) subjecting said SAM to thermal treatment so as to change

5 the energy-balance of said SAM in the thermally treated area so that said SAM is converted into a different structural form, preferably a more stable form.

10 In a preferred form of the invention, there is used a thermal treatment applied selectively to only part of the SAM in accordance with a negative image of said desired pattern so that the thermally treated areas are rendered more stable compared to the untreated ones.

15 Thus by means of the present invention it is possible to provide to produce patterned SAMs in a novel manner which differs from previously known processes. Further surprising and unexpected features of the invention include inter alia, the significantly increased structural perfection of the thermally treated SAM
20 with significantly increased domain size and resistance to structural disruption by surface discontinuities in the substrate surface.

The present invention can be used with a wide range of
25 substrates and compounds. Suitable substrates generally comprise conductors or semiconductors such as gold, silver,

chromium, manganese, vanadium, tungsten, molybdenum, zirconium, titanium, platinum, aluminium, iron, steel, indiumphosphide, gallium arsenide, and alloys and oxides, including glasses such as silicates and borates, as well as mixtures of such materials.

5

Suitable compounds generally comprise an anchor moiety, and an optionally substituted, aryl (including hetero-aryl) moieties, and optionally a spacer moiety between the anchor moiety and the aryl moiety. Suitable anchor moieties include thio, seleno, 10 carboxyl, phosphonate, phosphate and hydroxyl. Suitable aryl moieties include phenyl, biphenyl and terphenyl, as well as fused ring systems such as anthracyl and naphthyl, and hetero-aryl groups such as bipyridyl, terpyridyl, thiophenyl, bithienyl, terthienyl and pyrrolyl. Optional aryl substituents 15 include halogen, carboxy, trifluoromethyl, thiol, hydroxy, cyano, amino, nitro, lower alkyl e.g. C1 to C6 and carbonyl. The use of such substituents can be useful in modifying the properties of the SAM in generally known manner. Thus, for example, non-polar substituents such as CH₃ can be used to make 20 the SAM surface more hydrophobic, and polar substituents such as OH or COOH can be used to make the SAM surface more hydrophilic.

Suitable spacer groups include low molecular weight, saturated or unsaturated hydrocarbon chains and/or other structures 25 containing e.g. ether linkages, amide groups, or even e.g. cycloalkyl cyclic structures. The spacer groups are preferably

C1 to C10, advantageously C1 to C10 alkyl, most preferably C1 to C6. As further discussed hereinbelow, a number of factors affects the energy balance of the SAM and it is necessary that at least two of these factors enter the energy balance in a competitive way. There is a directive force originating from the SAM molecule-substrate interface, i.e. the substrate-head group bonding geometry favours a certain orientation of the molecule. (For the avoidance of doubt, it should be noted that the term "head group" is used herein to indicate the anchor moiety of the SAM molecule which is bound to the substrate, the other end being referred to as the "tail group"). A further important factor is the intermolecular interaction between neighbouring SAM molecules (?) which is maximized in particular orientations of the SAM molecules. Design of the molecular structure is generally effected in such a way that the two competing factors cannot be maximized at the same time whereby the resulting SAM film structure represents a compromise between the competing factors which, therefore, results in pronounced local minima on the energy hypersurface of the system. As a consequence structural transitions between different (meta)stable structures and associated changes in properties are possible.

Additional factors entering the energy balance to a greater or lesser extent are the strength of the head group-substrate interaction, the head group-substrate corrugation potential (variation of the SAM molecule - substrate interaction across the substrate surface), possible reconstructions of the

substrate surface at the head group-substrate interface, interactions between the SAM and its environment, and/or conformational degrees of freedom of the adsorbed molecules.

- 5 The molecular structure generally should match the substrate, e.g. for thiols on gold an even number of carbon atoms in the hydrocarbon spacer group between the head group and the aryl moiety, is required (C2, C4, C6, C8, C10) due to an sp^3 -like bonding geometry of the substrate-sulphur-carbon bond angle
- 10 whereas on silver an odd number (C1, C3, C5, C7, C9) produces the corresponding structure due to an sp^3 -like bonding geometry of the substrate-sulphur-carbon bond angle. Care should be taken to avoid situations in which one of the factors dominates too strongly, e.g. if the intermolecular interactions dominates
- 15 excessively the substrate-head group bonding geometry, then the SAM will not be able to transform under thermal treatment in accordance with the present invention. Thus, if for example, in the case of SAM molecules comprising unsubstituted biphenyl moieties, the length of the alkane spacer group might be limited
- 20 to less than ten carbon atoms since otherwise the intermolecular interactions start to dominate excessively the substrate-head group bonding geometry.

One preferred group of compounds suitable for use as the SAM

25 compounds of the present invention are 4-(4'-Methyl-biphenyl-4-yl)-alkane-1-thiols, especially the C1 to C10 (alkane)

compounds. These are conveniently referred to as BPn where n is the number of carbon atoms in the alkane moiety, thus BP4 corresponds to 4-(4'-Methyl-biphenyl-4-yl)-butane-1-thiol.

5 The SAMs obtained using such selected compounds may be heat treated in various different ways. Thus they may be heated by means of direct contact with a heated body including thermal nanolithography with a microscopic heated tip tool), or a heated fluid (liquid or gas). Thermal treatment may also be effected
10 remotely by means of thermal radiation including infra-red radiation and laser radiation, which are generally easier to control. Such radiation may be continuous or pulsed, the latter being preferred in order to avoid loss of resolution in the image transfer process due to heating of areas adjacent to those
15 being heated directly, as a result of thermal conduction from the latter to the former, resulting in unintentional heating of the former. Pulsed laser and other radiation can be particularly convenient in view of the various different irradiation parameters (pulse profile including height (energy)
20 and duration, as well as duration of inter-pulse interval) which can be more or less readily controlled.

It will be appreciated that the treatment temperatures required for different SAMs and/or different SAM-substrate combinations,
25 may differ to some extent. In general, though, we have found that the SAM should be raised to a temperature of at least

around 100 to 140°C. On the other hand excessively high temperatures should be avoided as these may result in disruption of the physical structure, and/or chemical degradation of the SAM compounds, or simply formation of yet other structural forms which are undesirable, e.g. because they are less stable and/or less resistant to exchange. In general, though, the treatment conditions used will involve a balance between factors such as temperature, treatment time, treatment mode (e.g. pulsed or continuous irradiation), and desired resolution).

10

In order to protect the SAM against chemical degradation and/or contamination it is desirable, to use a substantially inert atmosphere. Thus the heat treatment may be carried out under vacuum, or a noble gas, such as Argon or a relatively inert gas such as nitrogen.

15

The duration of the thermal treatment (not including the interruptions in the case of pulsed mode treatments), can also be varied and will moreover depend to a greater or lesser extent on the treatment temperature used. In general, higher temperatures reduce the time of treatment and typical values for a biphenyl thiol SAM are 150°C and 15 hrs for continuous (non-pulsed) treatment. Insofar as the change in crystalline structure and/or packing density can be readily monitored by means of scanning probe microscopies, ellipsometry, vibrational spectroscopy (applicable in situ), and/or ex situ by

20

25

-11-

structurally sensitive surface analysis methods or contact angle measurement (the latter after a suitable development process), it will be appreciated that suitable thermal treatment times for any given case can be readily determined by trial and error. By way of illustration we have found that the required structural form change can be achieved for a BP4 (4-(4'-Methyl-biphenyl-4-yl)-butane-1-thiol) SAM on gold using continuous thermal treatment of several hours, and with laser irradiation shorter than 1 hour, without significant degradation or chemical change in the SAM.

As noted above, various kinds of thermal treatment may be used in accordance with the present invention. Besides non-patterned treatment (where the whole SAM is treated uniformly), patterns can be generated by serial and parallel methods. Suitable types of treatment include contact methods (oven/hot plate for uniform treatment, and heated mask or tip in thermal contact with the SAM for patterned treatment) and contactless treatment (radiation, laser etc). For parallel processing (where the desired image is transferred simultaneously by exposure of the SAM through a suitable mask), a laser or any other light source of sufficient power is preferably used a principal advantage of parallel processing (irradiation through a mask or a mask in thermal contact with sample) is the speed with which the image can be transferred.

In some cases, though, where it is desired to increase resolution, serial processing (where the different parts of the image are transferred successively by "writing" them with a scanning beam or a point probe in thermal contact with the
5 sample), may be preferred. In this case there may be used a focused laser beam, a scanning near field optical tip or a heated tip. In this case it will be appreciated that the scanning speed may be used to control the degree of thermal treatment applied to the SAM. Moreover, by varying the scanning
10 speed, different parts of the image being transferred, may be subjected to different levels of thermal energy in a relatively simple and easily controlled manner.

Once an image has been thermally transferred to the SAM, it may
15 be "fixed" or "developed", by making use of the difference in properties between the original, untreated, SAM structure, and the new SAM structure produced by the thermal treatment. A principal difference is that the thermally treated structure is more resistant to exchange of the anchored compounds of the SAM
20 with other molecules, e.g. SAM forming thiols of various lengths, typically C4 to C20, for example, ω -mercaptohexadecanoic acid (MHA) or ω -mercaptoundecanoic acid). Such exchange will result in removal of the non-thermally treated, less stable structural form, and formation of an
25 adsorbate layer of the respective molecule. In general such replacement is carried out in solution, i.e. the thermally

treated SAM is exposed to a solution which contains the displacing molecules.

The patterned SAMs provided by the present invention may be used
5 for various different purposes involving greater or lesser degrees of further processing. The patterned SAMs will generally be developed or fixed in some way, for example, by using differences in the structure to treat the underlying substrate (e.g. wet chemical etching) or to control
10 electrochemical processes, by modification of the SAM itself (e.g. particle or photon irradiation,) or by displacement of the less stable parts via exchange with other molecules. Such developed/fixed patterned SAMs can be used directly for, inter alia, controlling wetting properties, electrode properties
15 (conducting vs insulating, spatially defined change of work function), electrochemical, and/or tribological properties.

The patterned SAMs can also be used in conjunction with further processing of the substrate and/or to build additional material
20 layers etc. Thus, for example, the patterned SAM can be used as a lithographic mask for processing of the substrate, for example, by chemical and/or physical etching of those parts of the substrate with non-thermally treated SAM. Furthermore, the "patterned" (without further processing) or "fixed" (after
25 exchange of non treated areas by other molecules or other treatment) SAM can be used as template to direct electrochemical

processes (e.g. metal deposition by using a combination of conductive and blocking molecules) or chemical reactions (e.g. selective grafting of other molecules by using a combination of molecules bearing chemically active/passive end groups).

5

Further preferred features and advantages of the invention will appear from the following Examples and Figures provided by way of illustration. In the Figures:

Fig. 1 is a schematic cross-sectional view showing the effect of the spacer chain length on the arrangement of the anchored compound in the SAM;

Fig. 2 is a schematic illustration of a thermal treatment processing apparatus;

Fig. 3 is a view corresponding to Fig. 1b showing change of some of the anchored compound following thermal treatment;

Fig. 4 shows STM and optical microscopy images of different structural forms of BP4 SAMs;

Fig. 5 shows a graph of the difference in change of contact angle following exchange treatment with MHA of different BP4 SAM structural forms;

Fig. 6 is a view corresponding to Fig. 3 showing "fixing" of the patterned SAM of Fig. 3 by exchange treatment; and

Fig. 7A and 7B illustrate schematically, different patterning arrangements.

25

Example 1 - Preparation of Patterned SAM

A - Preparation of SAM

BP4 (4-(4'-Methyl-biphenyl-4-yl)-butane-1-thiol) was prepared as described in Rong et al, Langmuir 17, 1582 (2001). A BP4 SAM was then prepared at room temperature by immersion of a gold
5 substrate (polycrystalline gold (111) film (100nm thick evaporated at 2nm/s onto mica at 340°C and flame amended in an oxygen flame) into a solution of BP4 (10 micromolar) in ethanol for typically 24h. Subsequently, the coated substrate specimen was rinsed with ethanol and blown dry with nitrogen or argon.
10 Fig. 1(b) shows schematically the BP4 SAM obtained and Fig. 1 (a) shows for comparison an analogous BP3 SAM.

B - Thermal Treatment of SAM

The SAM was heated in a closed container (as illustrated in Fig.
15 2) which was filled with nitrogen at 150°C for 15 hrs.

C - Structural Properties of Patterned SAM

Thermally treated and non-treated areas of the BP4 SAM differ both in molecular density and structural perfection. The non-
20 treated structure adopts a $(5\sqrt{3} \times 3)$ structure with an area of 27 Å² per molecule (α . phase) whereas the annealed structure adopts a $(6\sqrt{3} \times 2\sqrt{3})$ structure and an area of 32.4 Å² per molecule (β . phase). With domain sizes of the β -phase easily exceeding 10⁵ nm² compared to typically < 10³ nm² of the α -phase the structural
25 perfections of the SAM is dramatically improved upon annealing.

The thermally treated and non-treated BP4 SAM areas are also compared schematically in Fig. 3.

Fig. 4A-D shows scanning tunneling microscope images of BP4 SAM on a gold substrate: (A) is a large scale image showing phase α (bright areas) and β phase (darker areas) coexisting. (B), (C) are more detailed molecular resolved images showing molecular packing and illustration of the unit cell. Each spot represent a molecule. Differences in brightness correspond to differences in tunneling current. The sample shown in (A) displays random patterning obtained by incomplete thermal treatment (at a sub-optimal treatment temperature) whereas the sample shown in (D) shows complete transformation from α to β phase. White arrows indicate domain boundaries, and black arrows indicate monoatomic steps in the gold substrate.

Fig. 4E is an optical microscope image, showing a condensation pattern on top of a BP4 SAM on gold, in which a pattern has been "developed" by immersion in mercaptohexadecanoic acid (MHA). Since MHA is significantly more hydrophilic than BP4, water condenses preferentially on those areas (the hexagons) occupied by MHA in place of the BP4 molecules. The size of the hexagons is about 60 μm .

Example 2 - "Fixing" of Patterned SAM

-17-

The thermally treated SAM obtained in Example 1, was immersed in a 1mM solution of ω -mercaptohexadecanoic acid (MHA) in ethanol for differed periods of time (from 5 minutes to more than 30 days) at ambient temperature. As may be seen in the upper curve
5 in Fig. 5, even after 4 hours treatment with MHA there is little more than 5% change in contact time, indicating very little exchange of the SAM. In contrast, treatment of a corresponding non-thermally treated SAM with MHA, results in a large change of some 40% in contact angle indicating a high rate of exchange of
10 SAM compound with MHA. Fig. 6 illustrates schematically the replacement of thermally non-treated BP4 SAM areas with MHA.

Fig.7A shows a SAM layer 1 on a gold substrate 2, being selectively thermally treated by irradiation with laser
15 radiation 3 though a mask 4. Fig.7B shows shows a SAM layer 1 on a gold substrate 2, being selectively thermally treated by means of a 2D scanning microscopic heated tip tool 5.



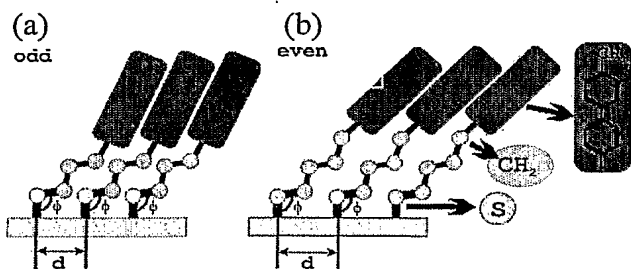


Fig. 1

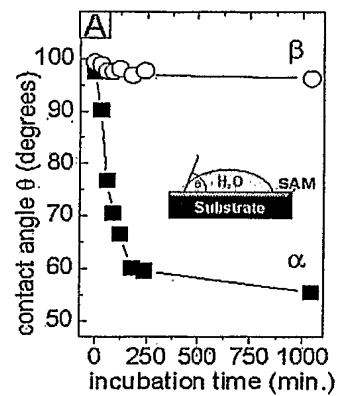


Fig. 5

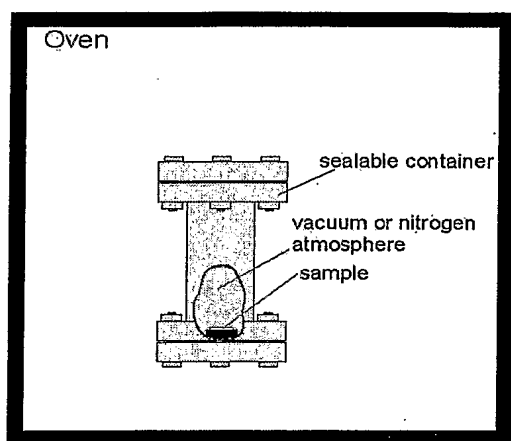


Fig. 2

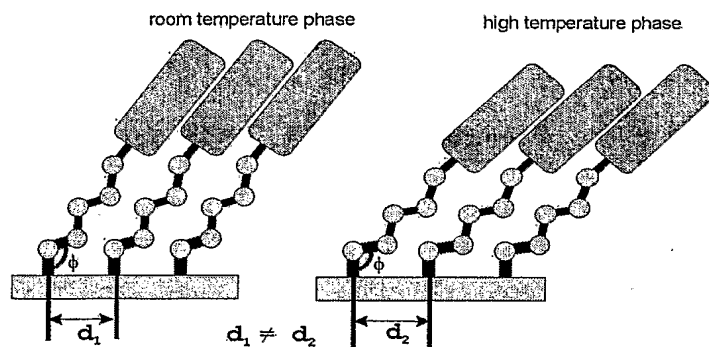


Fig. 3

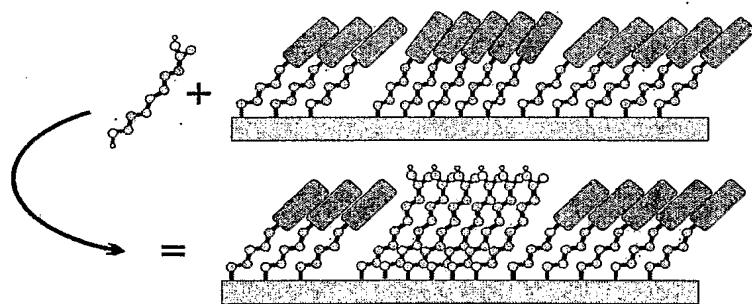


Fig. 6



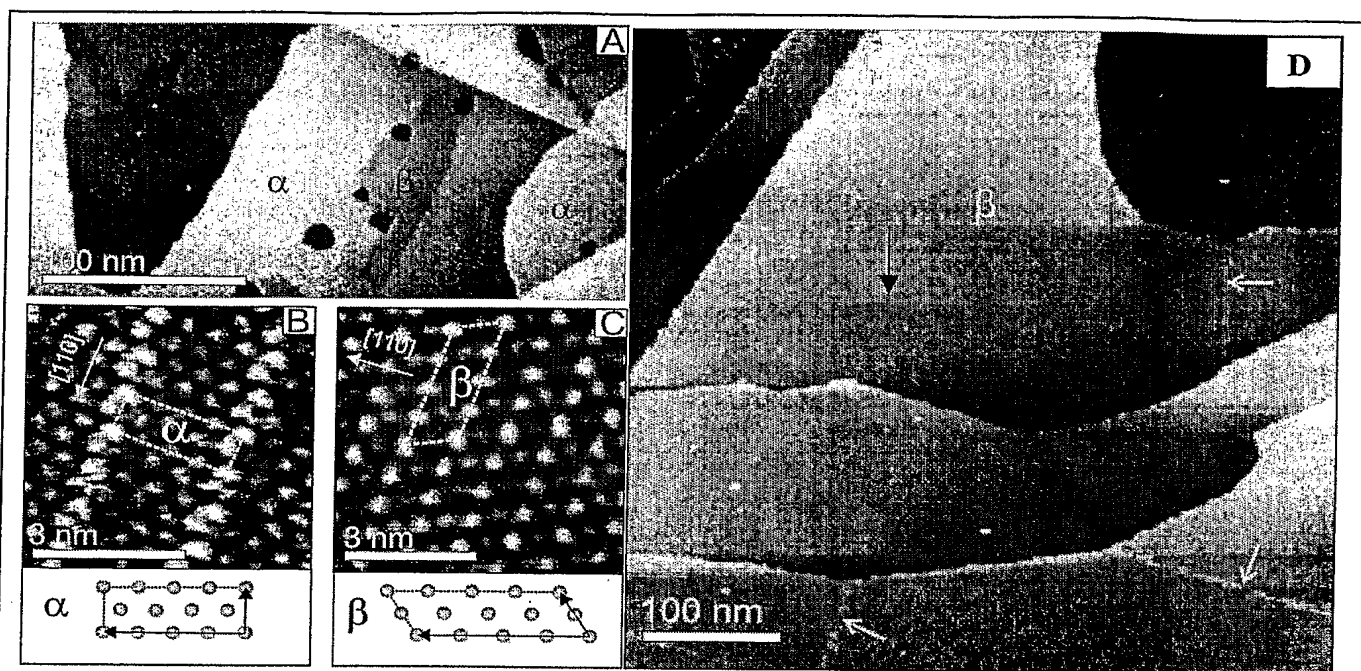
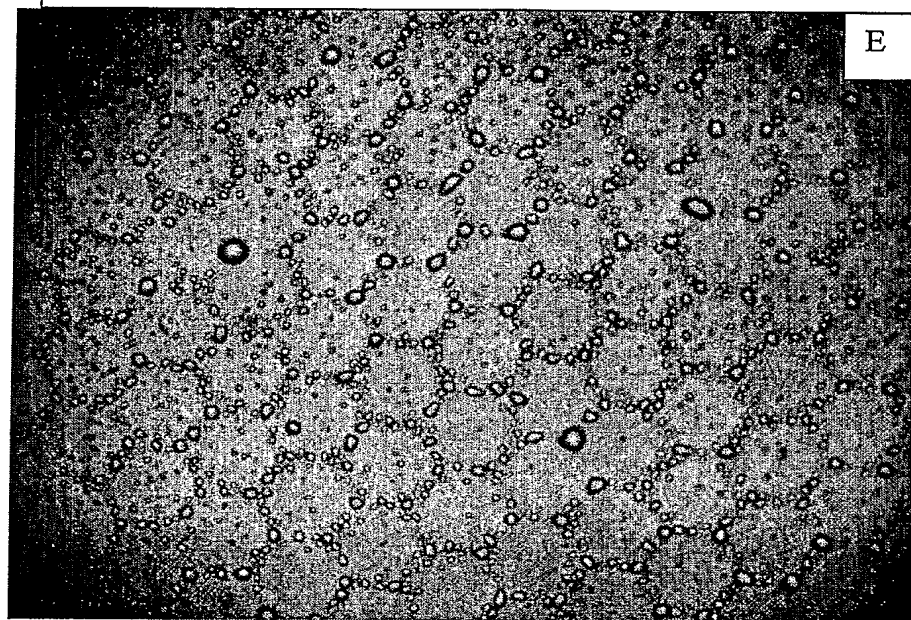
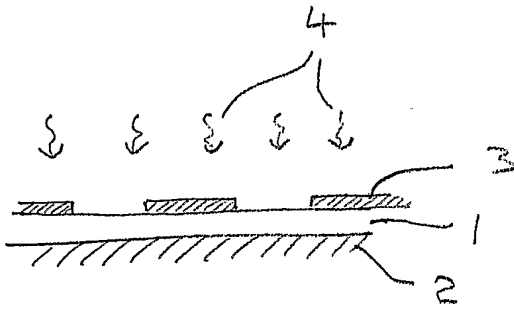
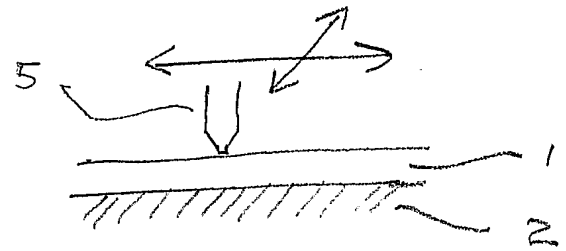


Fig. 4





FIG. 7AFIG. 7B

